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(54) **PURIFIED IgG ANTIBODIES**

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ANTICORPS IgG PURIFIES

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EP 0 504 363 B1



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L2: Entry 1 of 1

File: DWPI

Jul 29, 2003

DERWENT-ACC-NO: 1992-167163

DERWENT-WEEK: 200356

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TITLE: Immunosuppressant purified anti-CDW52 antibody preparations - for treating e.g. T-cell mediated diseases, rheumatoid arthritis, multiple sclerosis, diabetes, asthma, cancer, etc.

INVENTOR: ALLEN, G; RAMAGE, P I N ; RAMAGE, P I ; RAMAGE, P I M ; RAMAGAGE, P I N

PATENT-ASSIGNEE: WELLCOME FOUND LTD (WELL), BURROUGHS WELLCOME CO (WELL)

PRIORITY-DATA: 1990GB-0022547 (October 17, 1990)

Search Selected

Search ALL

Clear

PATENT-FAMILY:

| | PUB-NO | PUB-DATE | LANGUAGE | PAGES | MAIN-IPC |
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DESIGNATED-STATES: AT AU CA CH ES FI GB HU JP KR LU US AT BE CH DE DK ES FR GB GR IT LU NL SE
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APPLICATION-DATA:

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ABSTRACTED-PUB-NO: EP 504363B

BASIC-ABSTRACT:

Following are described. (A) purified prepn. of an anti-CDw52 antibody which exhibits on size exclusion chromatography a single peak under non-reducing conditions and 2 major peaks under reducing conditions; esp. the prepn. exhibits on conventional SDS polyacrylamide gel electrophoresis, one main band using a non-reduced sample and two main bands using a reduced sample; (B) purified prepn. of an anti-CDw52 antibody having a specific activity greater than 0.8 KU/mg; (C) purified prepn. of an anti-CDw52 antibody, free from host cell contaminants and/or aggregates; (D) purifying an anti-CDw52 antibody comprising (a) applying an aqs. soln. of the antibody to a Protein A column so as to absorb the antibody and eluting the antibody with an acid soln., (b) applying the acidic eluate to an ion-exchange column of charged particles to absorb the antibody and then eluting the antibody with an aqs. soln. of counter-charged ions and (c) applying the aqs. eluate to a size exclusion column of porous particles to retain non-antibody mols. in the porous particles and to obtain the desired antibody in selected fractions produced from the column.

Pref. the Protein A column is Protein A Sepharose and the antibody is eluted with citric acid.

USE - Purified anti-CDw52 antibodies are useful as immunosuppressives for treating T-cell mediated disorders including severe vasculitis, rheumatoid arthritis, systemic lupus, and also

autoimmune disorders such as multiple sclerosis, graft vs. host disease, psoriasis, juvenile onset diabetes, Sjogren's disease, thyroid disease, myasthenia gravis, transplant rejection and asthma. Antibodies are also useful in treating cancers e.g. Non-Hodgkins lymphoma and leukemias.

ABSTRACTED-PUB-NO: US 5644036A
EQUIVALENT-ABSTRACTS:

Following are described. (A) purified prepn. of an anti-CDw52 antibody which exhibits on size exclusion chromatography a single peak under non-reducing conditions and 2 major peaks under reducing conditions; esp. the prepn. exhibits on conventional SDS polyacrylamide gel electrophoresis, one main band using a non-reduced sample and two main bands using a reduced sample; (B) purified prepn. of an anti-CDw52 antibody having a specific activity greater than 0.8 KU/mg; (C) purified prepn. of an anti-CDw52 antibody, free from host cell contaminants and/or aggregates; (D) purifying an anti-CDw52 antibody comprising (a) applying an aqs. soln. of the antibody to a Protein A column so as to absorb the antibody and eluting the antibody with an acid soln., (b) applying the acidic eluate to an ion-exchange column of charged particles to absorb the antibody and then eluting the antibody with an aqs. soln. of counter-charged ions and (c) applying the aqs. eluate to a size exclusion column of porous particles to retain non-antibody mols. in the porous particles and to obtain the desired antibody in selected fractions produced from the column.

Pref. the Protein A column is Protein A Sepharose and the antibody is eluted with citric acid.

USE - Purified anti-CDw52 antibodies are useful as immunosuppressives for treating T-cell mediated disorders including severe vasculitis, rheumatoid arthritis, systemic lupus, and also autoimmune disorders such as multiple sclerosis, graft vs. host disease, psoriasis, juvenile onset diabetes, Sjogren's disease, thyroid disease, myasthenia gravis, transplant rejection and asthma. Antibodies are also useful in treating cancers e.g. Non-Hodgkins lymphoma and leukemias.

Process for obtaining a purified IgG antibody preparation which comprises:

(a) culturing a recombinant mammalian cell line capable of producing the antibody in an aqueous culture medium under antibody producing conditions;

(b) applying the antibody-containing aqueous medium to a Protein A column or Protein G column so as to absorb the antibody onto the column and then eluting the antibody with an acidic solution to produce an acidic eluate;

(c) applying the acidic eluate to an ion exchange column of charged particles previously equilibrated with a neutral buffer so as to absorb the antibody and then eluting the antibody with a neutral aqueous solution of counter-charged ions; and

(d) applying the aqueous eluate to a size exclusion column of porous particles so as to separate aggregates and other non-antibody molecules from the desired antibody and to obtain the desired antibody in selected fractions eluted from the column.

WO 9207084A

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DERWENT-CLASS: B04 D16

CPI-CODES: B04-B04C; B12-A07; B12-C10; B12-D02; B12-D03; B12-D07; B12-E01; B12-E02; B12-G05; B12-G06; B12-G07; B12-K02; D05-H11;

ANSWER 3 OF 3 WPIDS COPYRIGHT 2004 THOMSON DERWENT on STN

AN 1992-167163 [20] WPIDS

DNC C1992-076899

TI Immunosuppressant purified anti-CDW52 antibody preparations -
for treating e.g. T-cell mediated diseases, rheumatoid arthritis,
multiple sclerosis, diabetes, asthma, cancer, etc..

DC B04 D16

IN ALLEN, G; RAMAGE, P I N; RAMAGE, P I; RAMAGE, P I M; RAMAGAGE, P I N

PA (WELL) WELLCOME FOUND LTD; (WELL) BURROUGHS WELLCOME CO

CYC 28

PI WO 9207084 A1 19920430 (199220)* EN 43p

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Following are described. (A) purified preparation of an anti-CDw52 antibody which exhibits on size exclusion chromatography a single peak under non-reducing conditions and 2 major peaks under reducing conditions; especially the preparation exhibits on conventional SDS polyacrylamide gel electrophoresis, one main band using a non-reduced sample and two main bands using a reduced sample; (B) purified preparation of an anti-CDw52 antibody having a specific activity greater than 0.8 KU/mg; (C) purified preparation of an anti-CDw52 antibody, free from host cell contaminants and/or aggregates; (D) purifying an anti-CDw52 antibody comprising (a) applying an aqs. solution of the antibody to a Protein A column so as to absorb the antibody and eluting the antibody with an acid solution, (b) applying the acidic eluate to an ion-exchange column of charged particles to absorb the antibody and then eluting the antibody with an aqs. solution of counter-charged ions and (c) applying the aqs. eluate to a size exclusion column of porous particles to retain non-antibody mols. in the porous particles and to obtain the desired antibody in selected fractions produced from the column.

Pref. the Protein A column is Protein A Sepharose and the antibody is eluted with citric acid.

USE - Purified anti-CDw52 antibodies are useful as immunosuppressives for treating T-cell mediated disorders including severe

vasculitis, rheumatoid arthritis, systemic lupus, and also autoimmune disorders such as **multiple sclerosis**, graft vs. host disease, psoriasis, juvenile onset diabetes, Sjogren's disease, thyroid disease, myasthenia gravis, transplant rejection and asthma. Antibodies are also useful in treating cancers e.g. Non-Hodgkins lymphoma and leukemias. (0/0)
0/0

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